IN THE CLAIMS

Cancel claims 1-7.

Add the following claims:

8. Method for treating excessive blood lipid levels in humans, exclusively, with non-prescription, over-the-counter, food supplements consisting of:

formulating a combination of two 500 mg soft gel capsules of inositol hexanicotinate; two 1250 mg soft gel capsules, each containing about 450 mg eicosapentaenoic acid and about 200 mg docosahexaenoic acid; and two 1200 mg soft gel capsules, each containing about 192 mg phosphatidyl choline, about 168 mg phosphatidyl ethanolamine, and about 108 mg phosphatidyl inositol;

administering said combination orally, daily.

9. The method of claim 8 wherein said combination is administered for at least about six weeks.

8. (pending) Method for treating excessive blood lipid levels in humans, exclusively,

with non-prescription, over-the-counter, food supplements consisting of:

formulating a combination of two 500 mg soft gel capsules of inositol hexanicotinate; two 1250 mg soft gel capsules, each containing about 450 mg eicosapentaenoic acid and about 200 mg docosahexaenoic acid; and two 1200 mg soft gel capsules, each containing about 192 mg phosphatidyl choline, about 168 mg phosphatidyl ethanolamine, and about 108 mg phosphatidyl inositol;

administering said combination orally, daily.

9. (pending) The method of claim 8 wherein said combination is administered for at least about six weeks.

Pursuant to the requirements of 37 CFR Section 1.173 (c) the above describes the status of the two claims added in this case, i.e., pending. Support therefor is found in the specification as follows:

Support for the "over the counter" requirement is found for example at page 3, lines 16-18 and under the heading "Field of the Invention".

Support for using flush free niacin is found for example at page 5, lines 6-14.

Support for the remaining chemicals in claim 8 is found in the instant specification at page 7.